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=> file medline biosis caplus embase
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=> s nucleic (p)duplex(p)temperature L1 211 NUCLEIC (P) DUPLEX(P) TEMPERATURE

=> s l1(p)below(p)melting L2 6 L1(P) BELOW(P) MELTING

=> duplicate remove 12

DUPLICATE PREFERENCE IS 'MEDLINE, BIOSIS, EMBASE'

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PROCESSING COMPLETED FOR L2

L3 2 DUPLICATE REMOVE L2 (4 DUPLICATES REMOVED)

=> d his

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FILE 'MEDLINE, BIOSIS, CAPLUS, EMBASE' ENTERED AT 12:22:26 ON 09 JAN 2003 L1 211 S NUCLEIC (P) DUPLEX (P) TEMPERATURE

L2 6 S L1(P)BELOW(P)MELTING

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Main Mer	nu Search Form Posting Counts Show S Numbers Edit S Numbers Preferences Cases		
	Search Results - Terms Documents L2 same 35 2		
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DATE: Thursday, January 09, 2003 Printable Copy Create Case

Set Name Query side by side		Hit Count	Set Name result set	
DB=USPT; $PLUR=YES$; $OP=OR$				
<u>L6</u>	L2 same 35	2	<u>L6</u>	
<u>L5</u>	L2 same 48	1	<u>L5</u>	
<u>L4</u>	L2 same 40	0	<u>L4</u>	
<u>L3</u>	L2 same 40 same 48	0	<u>L3</u>	
<u>L2</u>	L1 same below same melting	140	<u>L2</u>	
<u>L1</u>	nucleic same duplex same temperature	1466	<u>L1</u>	

END OF SEARCH HISTORY

L3ANSWER 1 OF 2 MEDLINE DUPLICATE 1 97465962 AN MEDLINE DN 97465962 PubMed ID: 9321670 ΤI A fiber optic biosensor for fluorimetric detection of triple-helical DNA. ΑU Uddin A H; Piunno P A; Hudson R H; Damha M J; Krull U J CS Department of Chemistry, Otto Maas Chemistry Building, McGill University, Montreal, Quebec H3A 2K6, Canada. SO NUCLEIC ACIDS RESEARCH, (1997 Oct 15) 25 (20) 4139-46. Journal code: 0411011. ISSN: 0305-1048. CYENGLAND: United Kingdom DTJournal; Article; (JOURNAL ARTICLE) LΑ English FS Priority Journals EΜ 199712 ED Entered STN: 19980109 Last Updated on STN: 19990129 Entered Medline: 19971202 AB A fiber optic biosensor was used for the fluorimetric detection of T/AT triple-helical DNA formation. The surfaces of two sets of fused silica optical fibers were functionalized with hexaethylene oxide linkers from which decaadenylic acid oligonucleotides were grown in the 3'to 5'and 5'to 3'direction, respectively, using a DNA synthesizer. Fluorescence studies of hybridization showed unequivocal hybridization between oligomers immobilized on the fibers and complementary oligonucleotides from the solution phase, as detected by fluorescence from intercalated ethidium bromide. The complementary oligonucleotide, dT10, which was expected to Watson-Crick hybridize upon cooling the system below the duplex melting temperature (T m), provided a fluorescence intensity with a negative temperature coefficient. Upon further cooling, to the point where the pyrimidine motif T*AT triple-helix formation occurred, a fluorescence intensity change with a positive temperature coefficient was observed. The reverse-Hoogsteen T.AT triplex, which is known to form with branched nucleic acids, provided a corresponding decrease in fluorescence intensity with decreasing temperature. Full analytical signal evolution was attainable in minutes. L3 ANSWER 2 OF 2 MEDLINE DUPLICATE 2 AN 92253408 MEDLINE DN 92253408 PubMed ID: 1579489 Properties of pseudouridine N1 imino protons located in the major groove TIof an A-form RNA duplex. Hall K B; McLaughlin L W Department of Biochemistry and Molecular Biophysics, Washington University CS School of Medicine, St Louis, MO 63110. NC GM37065 (NIGMS) SO NUCLEIC ACIDS RESEARCH, (1992 Apr 25) 20 (8) 1883-9. Journal code: 0411011. ISSN: 0305-1048. ENGLAND: United Kingdom CY DT Journal; Article; (JOURNAL ARTICLE) LΑ English FS Priority Journals EM 199206 ED Entered STN: 19920619 Last Updated on STN: 19970203 Entered Medline: 19920605 AB The exchangeable N1 imino protons of two pseudouridine (psi) bases located at adjacent internal positions within an undecamer RNA duplex (5'AUAC psi psi ACCUG/3'UAUGAAUGGUC) can report on the environment of the

The exchangeable N1 imino protons of two pseudouridine (psi) bases located at adjacent internal positions within an undecamer RNA duplex (5'AUAC psi psi ACCUG/3'UAUGAAUGGUC) can report on the environment of the major groove of an A-form double-stranded nucleic acid. The psi N1 imino protons of these residues (which are not involved in interstrand Watson-Crick hydrogen bonding) are protected from chemical exchange with the solvent water and thus are observable in the proton NMR spectrum in

H2O (1). These protons will exchange readily at increased pH values or upon thermal denaturation of the duplex. The longitudinal (T1) relaxation times of the psi N1 imino protons in 100 mM NaCl or in 10 mM MgCl2 and 100 mM NaCl are approximately two-fold faster than those of the psi N3 imino protons which are involved in Watson-Crick base pairing. With the addition of spermidine, the psi N1 imino protons become readily exchangeable at a temperature some 20 degrees C below the melting temperature of the duplex.